B. SOLID-ROCK COLUMNS

Direct measurements of transport parameters in actual subsurface materials under subsurface conditions are necessary for defensible modeling of contaminant transport in host rocks and engineered barriers surrounding nuclear and hazardous waste repositories. The hydraulic conductivity, K, and the retardation factor, $R_{\rm f}$, along with the associated distribution coefficient, K_d , are poorly known transport parameters for real systems but are key input parameters to existing and developing contaminant release models. We experimentally determined unsaturated R_f and K for core samples of Yucca Mountain vitric-member tuff and zeolitic nonwelded tuff from G-Tunnel, Bed 5, with respect to J-13 well water with a selenium concentration (as selenite) of 1.31 mg/l (ppm) at 23°C. Our intent was to demonstrate that a method in which flow is induced with an ultracentrifuge (the UFATM method) could rapidly and directly measure $R_{\rm f}$ and K in whole-rock tuff cores and then to compare these directly measured unsaturated R_f values with those calculated from K_d values obtained through traditional batch tests on the same materials.

Methodology

Retardation

Retardation factors can be determined in flow experiments where $R_{\rm f}$ for a particular species is the ratio of the solution velocity to the species velocity. The retardation factor for that species is given by:

$$R_{\rm f} = \frac{V_{\rm gw}}{V_{\rm sp}} = 1 + \rho_{\rm d} \frac{K_{\rm d}}{\epsilon} , \qquad (46)$$

where $V_{\rm gw}$ is the velocity of carrier fluid, $V_{\rm sp}$ is the velocity of the species, $\rho_{\rm d}$ is the dry bulk density, ϵ is the porosity, and $K_{\rm d}$ is defined as the moles of the species per g of solid divided by the moles of the species per ml of solution. If none of a particular species is lost to the solid phase, then $K_{\rm d}=0$ and $R_{\rm f}=1$ for that species. In column experiments, a breakthrough curve is obtained for the particular species and $R_{\rm f}$ is determined as the pore volume at which the concentration of the species in

the solution that has passed through the column is 50% of the initial concentration ($C/C_0=0.5$). It is now generally assumed that for unsaturated systems $\epsilon=\theta$, where θ is the volumetric water content (Bouwer 1991; Conca and Wright 1992a). The study described in this section experimentally addresses this concern under unsaturated conditions in whole rock and evaluates the use of data from batch experiments in determining $R_{\rm f}$ in whole rock.

We prepared solutions using J-13 well water with a selenite concentration of 1.31 ppm and determined the selenium concentrations with an inductively coupled, argon-plasma, atomic-emission spectrometer (Jarrell-Ash Model 976 Plasma Atomcomp). We used an ion chromatograph (Dionex Series 4000i) to determine the speciation of selenium in solution. All selenium in the starting and effluent solutions was found to exist as selenite.

Hydraulic conductivity

One way to drive fluid through rock is to use centripetal acceleration as the driving force. We used this approach with a new technology (UFA) to produce hydraulic steady-state, to control temperature, degree of saturation, and flow rates in all retardation experiments, and to measure the hydraulic conductivity. A specific advantage of this approach is that centripetal acceleration is a whole-body force similar to gravity that acts simultaneously over the entire system and independently of other driving forces, such as gravity or matrix suction. It has been shown that capillary bundle theory holds in the UFA method (Conca and Wright 1992a, 1992b).

The UFA instrument consists of an ultracentrifuge with a constant, ultralow flow-rate pump that provides fluid to the sample surface through a rotating seal assembly and microdispersal system (Fig. 116). Accelerations up to 20,000 g are attainable at temperatures from 220° to 150°C and flow rates as low as 0.001 ml/hr. The effluent is collected in a transparent, volumetrically calibrated container at the bottom of the sample assembly. The effluent

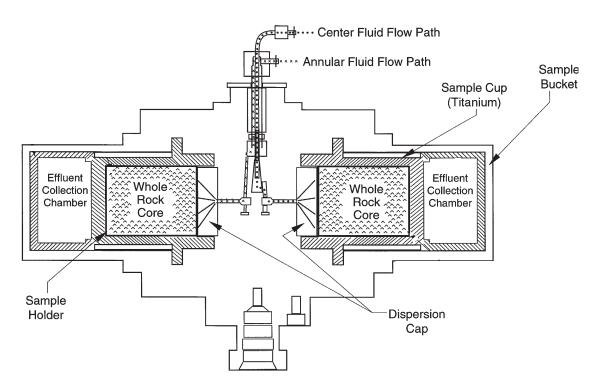


Figure 116. The UFA Method. This schematic of the UFA rotor is shown with paramagnetic seal and the large-sample options, a configuration that is optimal for adsorption and retardation studies.

collection chamber can be observed during centrifugation using a strobe light.

The current instrument has two different rotor sizes that hold up to 50 and 100 cm³ of sample, respectively. Three different rotating-seal assemblies facilitate various applications and contaminant compatibilities; they are a face seal, a mechanical seal, and a paramagnetic seal. Figure 116 shows the large-sample option with the paramagnetic seal, a configuration that is optimal for adsorption and retardation studies.

Numerous studies have compared use of the UFA approach with traditional methods of doing this type of analysis in soils and clays, and the agreement is excellent (Conca and Wright 1992b; Nimmo et al. 1987). Good agreement is expected because the choice of driving force does not matter provided the system is Darcian (see next paragraph) and the sample is not adversely affected by a moderately high driving force ($\leq 1000 \text{ g}$ for all

samples run in these experiments); both of these provisions hold for most geologic systems. Additionally, all techniques for estimating hydraulic conductivity, $K(\theta)$, are extremely sensitive to the choice of the rock or soil residual water content, $\theta_{\rm r}$, and to the saturated hydraulic conductivity, $K_{\rm s}$; minor variations in $\theta_{\rm r}$ or $K_{\rm s}$ produce order-of-magnitude changes in $K(\theta)$ (Stephens and Rehfeldt 1985).

The UFA technology is effective because it allows the operator to set the variables in Darcy's Law, which can then be used to determine hydraulic conductivity. Under a centripetal acceleration in which water is driven by both the potential gradient, $d\psi/dr$, and the centrifugal force per unit volume, $\rho\omega^2 r$, Darcy's Law is

$$q = -K(\psi)\left[\frac{d\psi}{dr} - \rho\omega^2 r\right] , \qquad (47)$$

where q is the flux density into the sample; K, the hydraulic conductivity, is a function of the matric

suction (ψ) and, therefore, of water content (θ) ; r is the radius from the axis of rotation; ρ is the fluid density; and ω is the rotation speed. When multicomponent and multiphase systems are present in the UFA instrument, each component reaches its own steady-state with respect to each phase, as occurs in the field. Appropriate values of rotation speed and flow rate into the sample are chosen to obtain desired values of flux density, water content, and hydraulic conductivity in the sample. Above speeds of about 300 rpm, depending upon the material and providing that sufficient flux density exists, $d\psi/dr \ll \rho\omega^2 r$. Under these conditions, Darcy's Law is given by $q = -K(\psi) \left[-\rho \omega^2 r\right]$. Rearranging the equation and expressing hydraulic conductivity as a function of water content, Darcy's Law becomes

$$K(\theta) = \frac{q}{\rho \omega^2 r} \ . \tag{48}$$

As an example, a whole-rock core of Topopah Spring Member tuff accelerated to 7500 rpm with a flow rate into the core of 2 ml/hr achieved hydraulic steady-state in 30 hours with a hydraulic conductivity of 8.3×10^{-9} cm/sec at a volumetric water content of 7.0%. Previous studies have verified the linear dependence of K on flux and the second-order dependence on rotation speed (Conca and Wright 1992a; Nimmo et al. 1987), and several comparisons between the UFA method and other techniques have shown excellent agreement (Conca and Wright 1992a, 1992b). Because the UFA method can directly and rapidly control the hydraulic conductivity, fluid content, temperature, and flow rates, other transport properties can then be measured as a function of fluid content by associated methods either inside or outside the UFA instrument during the overall run.

Fundamental physics issues involving flow in an acceleration field have been raised and successfully addressed by previous research and in numerous forums (Conca and Wright 1992a, 1992b; Nimmo et al. 1987; Nimmo and Akstin 1988; Nimmo and Mello 1991). These studies have shown, first, that compaction from acceleration is negligible for sub-

surface soils at or near their field densities. Bulk density in all samples remain constant because a whole-body acceleration does not produce high point pressures. A notable exception is surface soils, which can have unusually low bulk densities; special arrangements must be made to preserve their densities. Whole-rock cores are completely unaffected.

The studies have also shown that three-dimensional deviations of the driving force with position in the sample are less than a factor of 2, but moisture distribution is uniform to within 1% in homogeneous systems because water content depends only upon ψ , and unit gradient conditions are achieved in the UFA instrument in which $d\psi/dr = 0$. Hydraulic steady-state is not as sensitive to changes in rotation speed as to flux density. In heterogeneous samples or multicomponent systems such as rock, each component reaches its own hydraulic steady-state and water content, as occurs for such materials under natural conditions in the field. This last effect cannot be reproduced with pressure-driven techniques but only under a wholebody force field, such as with gravity columns or centrifugal methods. The ratio of flux to rotation speed is always kept high enough to maintain the condition of $d\psi/dr = 0$.

Results and Discussion for Vitric and Zeolitic Tuff

Column breakthrough test results

For these experiments, the rotation speed was set at 2,000 rpm with a flow rate into each sample of 0.2 ml/hr. The experiment was run for 9 days with an initial selenium concentration of 1.31 ppm. Figure 117 shows the breakthrough curves for selenite in the Yucca Mountain vitric member at 62.6% saturation and in the zeolitic nonwelded tuff at 52.8% saturation. The experiment was stopped before full breakthrough in the zeolitic nonwelded tuff, but the $C/C_{\rm o}=0.5$ point was reached. The retardation factor for each tuff sample is only 2.5, giving a $K_{\rm d}$ of 0.9 ml/g for the Yucca Mountain vitric-member tuff and 0.8 ml/g for the zeolitic nonwelded tuff.

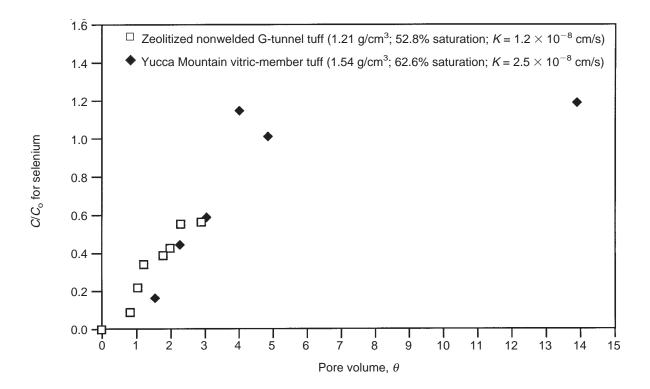


Figure 117. Breakthrough Curves. The UFA column data plotted here for a Yucca Mountain tuff retardation experiment show the breakthrough curves for selenium. The initial concentration, C_0 , of selenium (as selenite) was 1.31 ppm in J-13 well water.

During these experiments, the unsaturated hydraulic conductivity, K, for each sample at these water contents was 2.5×10^{-8} cm/s for the Yucca Mountain vitric-member tuff and 1.2×10^{-8} cm/s for the zeolitic nonwelded tuff. Figure 118 gives the characteristic curves, $K(\theta)$, for these tuffs determined in separate experiments, as well as measurements for other tuffs and materials for comparison. As in most whole-rock cores studied (Conca and Wright 1992a, 1992b), the characteristic curves for the tuffs are steep, almost linear functions of the volumetric water content and are displaced according to the degree of welding and alteration.

Batch test results

We conducted batch-adsorption tests using the same J-13 well water with the slightly lower selenium concentration of 1.1 ppm and the same zeolitic nonwelded tuff from G-Tunnel, Bed 5, as in the UFA column breakthrough test. The batch-adsorption tests consist of crushing and wet-siev-

ing the tuff, pretreating the tuff with J-13 water, placing the selenium solution in contact with the tuff, separating the phases by centrifugation, and determining the amount of selenium in each phase by difference using inductively coupled plasma mass spectrometry. Control samples were used to determine the sorption of selenium onto the walls of the sorption containers. The control procedure consisted of following the described batch-sorption procedure with a sample containing the selenium solution, except with no tuff added. The results of the control experiments indicate no loss of selenium due to precipitation or sorption onto the walls of the container during the batch-sorption experiment. The sorption distribution coefficients we obtained are given in Table 28. The Eh of all solutions, measured after the sorption experiments, varied from 140 to 150 mV.

The data presented in Table 28 and Fig. 118 indicate agreement between the column and the batch-

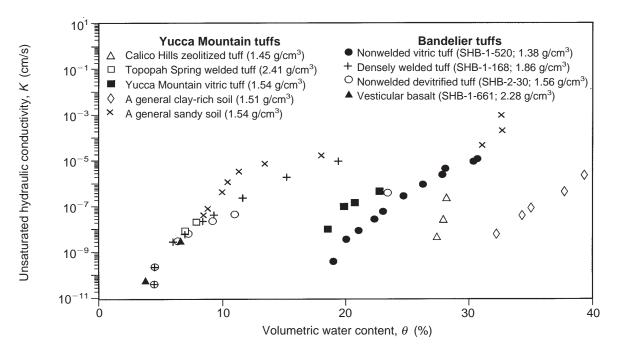


Figure 118. Unsaturated Hydraulic Conductivity. These UFA column data for various Yucca Mountain and Bandelier tuffs and other soil samples show the unsaturated hydraulic conductivity, K, as a function of volumetric water content, θ . The name and the density of each tuff is given in the legend.

sorption experiments. At a selenium concentration of ~1 ppm, no sorption of the selenium by the tuff is observed for the zeolitic tuff used in batch experiments, and minimal sorption (K_d of 0.8 ml/g) is observed for the zeolitic tuff used in the unsaturated column experiments. The method we used for

Table 28. Selenium Batch Adsorption on Nonwelded Zeolitic Tuff*

Pretreatment period (days)	Sorption period (days)	K _d (mL/g)
6.9	0.04	-0.2
6.9	0.04	0.3
6.8	13.9	0.0
6.8	13.9	0.2

^{*}Experimental conditions: J-13 water; 20° C; $75-500 \,\mu m$ tuff particle sizes; 1.1 ppm initial selenium concentration; solution pH after sorption of 8.4; and samples from the same location in G-Tunnel, Bed-5, as the tuff used in the column experiments.

the batch-sorption experiments to determine $K_{\rm d}$ values (by difference) involves subtracting the selenium concentration in solution after equilibration with the solid phase from the initial selenium concentration in solution. This method yields large scatter in the data when the batch-sorption distribution coefficient is small because two large numbers are subtracted to get a small number. Inspection of Table 28 also suggests that the kinetics of selenium sorption onto tuff are fast.

Conclusions

This study demonstrated the feasibility of using the UFA technology to rapidly and directly measure retardation factors and hydraulic conductivities in whole-rock cores of tuff under the unsaturated conditions that exist in the field. In UFA column breakthrough tests, the retardation factor for the selenite species was only 2.5 in both Yucca Mountain vitric member tuff at 62.6% saturation and zeolitic nonwelded tuff from G-tunnel at 52.8% saturation for a selenium concentration in

J-13 water of 1.31 ppm. In batch tests on the same material with an initial selenium concentration of 1.1 ppm, the average $K_{\rm d}$ was 0.08 ± 0.2 ml/g, which gives retardation factors that are slightly lower than those from the UFA column breakthrough experiments. This finding suggests that using batch-sorption coefficients to predict radionuclide transport through unsaturated tuff will yield conservative results.

Future experiments will use initial selenium concentrations smaller than the ones used in these experiments to further assess the validity of batch-sorption distribution coefficients to predict transport under unsaturated conditions. The unsaturated hydraulic conductivities during the experiments were 2.5×10^{-8} cm/s for the Yucca Mountain vitric-member tuff and 1.2×10^{-8} cm/s for the zeolitic nonwelded tuff.